PLOUG19.001APC PATENT

REQUEST FOR CORRECTED FILING RECEIPT

Applicant	:	Klaveness et al.
Appl. No.	:	10/583,829
Filed	:	April 5, 2007
For	:	MODULATORS OF PERIPHERAL 5-HT RECEPTORS
Art Unit	:	1614

Commissioner for Patents
P.O. Box 1450
Office of Initial Patent Examination
Customer Service Center
Alexandria, VA 22313-1450

Dear Sir:

Applicants hereby request that the Official Filing Receipt, a copy of which is enclosed, be corrected to reflect the true title of "MODULATORS OF PERIPHERAL 5-HT RECEPTORS". Presently, the Filing Receipt incorrectly shows the title as "MODULATORS OF PRERIPHERAL 5-HT RECEPTORS". The following is enclosed as evidence of the proper filing date:

(X) Copy of the first page of PCT/NO2004/000399.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: Sept 13, 2007

By: _____
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APPL NO.	FILING OR 371(c) DATE	ART UNIT	FIL FEE REC'D	ATTY.DOCKET NO	TOT CLMS	IND CLMS
10/583,829	04/05/2007	1614	1030	PLOUG19.001APC	19	1

CONFIRMATION NO. 8739

20995
KNOBBE MARTENS OLSON & BEAR LLP
2040 MAIN STREET
FOURTEENTH FLOOR
IRVINE, CA 92614

FILING RECEIPT

OC000000025043617

Date Mailed: 07/27/2007

Receipt is acknowledged of this nonprovisional patent application. The application will be taken up for examination in due course. Applicant will be notified as to the results of the examination. Any correspondence concerning the application must include the following identification information: the U.S. APPLICATION NUMBER, FILING DATE, NAME OF APPLICANT, and TITLE OF INVENTION. Fees transmitted by check or draft are subject to collection. Please verify the accuracy of the data presented on this receipt. **If an error is noted on this Filing Receipt, please write to the Office of Initial Patent Examination's Filing Receipt Corrections. Please provide a copy of this Filing Receipt with the changes noted thereon. If you received a "Notice to File Missing Parts" for this application, please submit any corrections to this Filing Receipt with your reply to the Notice. When the USPTO processes the reply to the Notice, the USPTO will generate another Filing Receipt incorporating the requested corrections (if appropriate).**

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Assignment For Published Patent Application

Bio-Medisinsk Innovasjon, Oslo, NORWAY

Power of Attorney: The patent practitioners associated with Customer Number **20995**.

Domestic Priority data as claimed by applicant

This application is a 371 of PCT/NO04/00399 12/23/2004

Foreign Applications

DENMARK PA 2003 01924 12/23/2003

If Required, Foreign Filing License Granted: 07/26/2007

The country code and number of your priority application, to be used for filing abroad under the Paris Convention, is US10/583,829

Projected Publication Date: 11/01/2007

Non-Publication Request: No

Early Publication Request: No

Title

Modulators of Preripheral 5-Ht Receptors

Preliminary Class

514

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(81) Designated States *(unless otherwise indicated, for every kind of national protection available)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

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Published:
— *without international search report and to be republished upon receipt of that report*

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WO 2005/061483 A2

(54) Title: MODULATORS OF PERIPHERAL 5-HT RECEPTORS

(57) Abstract: Novel modulators of 5-HT4 receptors have been developed which have a selectivity for peripheral receptors rather than those of the central nervous systems. Theses include novel derivatives of known modulators as well as entirely novel entities. Surprisingly, the derivatised compounds of the known modulators maintain a high binding affinity to 5-HT4 receptors, despite the presence of an acidic moiety at the end of an optional chain. The entirely novel entities also exhibit good binding affinity to 5-HT4 receptors. All of the compounds of the invention have a common motif which includes a basic nitrogen moiety and an acidic moiety. The compounds of the invention, due at least in part to their high ionisation potential at physiological pH, have the unique properties of selectively for peripheral 5HT4 receptors over those of the CNS, good binding affinity, and selectively of 5HT4 receptors over other serotonin receptors.